FDA 510(k) Premarket Notification

K761223 WIDEX HEARING AID MODEL F8

CLEARED

Dec 16, 1976
8 days to decision

K761223 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k761223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Dec 8, 1976
Decision date	Dec 16, 1976
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Widex Hearing Aid Co., Inc.**
Location	Mchenry, IL, US
510(k) history	52 submissions · 52 cleared · 1976-2008